UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/714,152	11/13/2003	Sekhar Boddupalli	0113-UTL2	2364
Gloria Pfister	7590 03/24/200	EXAMINER		
Galileo Pharma	The state of the s	QAZI, SABIHA NAIM		
	5301 Patrick Henry Dr. Santa Clara, CA 95054		ART UNIT	PAPER NUMBER
			MAIL DATE	DELIVERY MODE
			03/24/2009	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)
	10/714,152	BODDUPALLI ET AL.
Office Action Summary	Examiner	Art Unit
	Sabiha Qazi	1612
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim vill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).
Status		
Responsive to communication(s) filed on 18 No. This action is FINAL. 2b) ☑ This Since this application is in condition for allowant closed in accordance with the practice under E.	action is non-final. nce except for formal matters, pro	
Disposition of Claims		
4) ☐ Claim(s) 1-22 is/are pending in the application. 4a) Of the above claim(s) is/are withdraw 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 1-22 is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) are subject to restriction and/or Application Papers 9) ☐ The specification is objected to by the Examiner 10) ☐ The drawing(s) filed on is/are: a) ☐ acceptable.	vn from consideration. r election requirement. r. epted or b) □ objected to by the B	
Applicant may not request that any objection to the c Replacement drawing sheet(s) including the correcti	ion is required if the drawing(s) is obj	ected to. See 37 CFR 1.121(d).
11) The oath or declaration is objected to by the Ex	aminer. Note the attached Office	Action or form PTO-152.
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the prior application from the International Bureau * See the attached detailed Office action for a list of	s have been received. s have been received in Applicati ity documents have been receive ı (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 6/14./04.11/13/03.	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	nte

Application/Control Number: 10/714,152 Page 2

Art Unit: 1612

Non-Final Office Action

Claims 1-22 are pending. No claim is allowed at this time.

Summary of this Office Action dated 3/7/2009

- 1. Information Disclosure Statement
- 2. Copending Applications
- 3. Specification
- 4. 35 USC § 112 (1) Written Description Rejection
- 5. Response to Remarks
- 6. Communication

Examination of claims for purposes other than application of prior art, e.g. rejections under 35 USC 112, first and second paragraphs, will be understood to be made in the interest of completeness of prosecution and should not be considered as indicating that search has been extended to non-elected species.

Information Disclosure Statement

The listing of references in the specification is not a proper information disclosure statement. 37 CFR 1.98(b) requires a list of all patents, publications, or other information submitted for consideration by the Office, and MPEP § 609.04(a) states, "the list may not be incorporated into the specification but must be submitted in a separate paper." Therefore, unless the references have been cited by the examiner on form PTO-892, they have not been considered.

Copending Applications

Applicants must bring to the attention of the examiner, or other Office official involved with the examination of a particular application, information within their knowledge as to other copending United States applications, which are "material to patentability" of the application in question. MPEP 2001.06(b). See Dayco Products Inc. v. Total Containment Inc., 66 USPQ2d 1801 (CA FC 2003).

Specification

The specification has not been checked to the extent necessary to determine the presence of all possible minor errors. Applicant's cooperation is requested in correcting any errors of which applicant may become aware in the specification.

Claim Rejections - 35 USC § 112—Written Description Rejection

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 1-22 are rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement. The claim(s) contains subject matter, which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the

application was filed, had possession of the claimed invention. Following reasons apply. Following reasons apply:

Specification discloses in example s 1-5 use of oxazolone from Sigma chemical company. There is no data, teaching or guidance how to make and use the claimed invention. The specification does not contain the written description of the claimed subject matter.

Claims are drawn to a method of reducing the level of C-reactive protein (CRP) in an individual subject to a CRP associated inflammatory condition containing the compound of formula I. In formula I R can be O, S, SO, SO2, a secondary or tertiary amine, a phosphate, a phosphor-ester, or a substituted or unsubstituted methylene group. These compounds include thousands of compounds. See also when n can be 0 to 3, so that the hetero ring will vary depending on n.

Applicant had no possession at the time this application was filed of claimed subject matter. The test for determining compliance with the written description requirement is whether the disclosure of the application as originally filed reasonably conveys to one skilled in the art that the inventor had the possession at the time of the later claimed subject matter, rather than the presence or absence of literal support in the specification for the claimed language.

See In re Kaslow, 707 F 2d 1366, 1375 (Fed. Cir. 1983).

The description requirement of the patent statute requires a description of an invention, not an indication of a result that one might achieve if one made that invention. See, e.g., <u>In re Wilder</u>, 22 USPQ 369, 372-3 (Fed. Cir. 1984). (Holding that a claim was not adequately described because the specification did 'little more than outline goals appellants hope the claimed invention achieves and the problems the invention will hopefully ameliorate.')

Mere indistinct terms (such as "reducing the level of CRP and CRP associated condition" used herein), however, may not suffice to meet the written description requirement. This is particularly true when a compound is claimed in purely functional terms. See <u>Univ. of Rochester v. G.D. Searle</u>, 69 USPQ2d 1886 (CAFC 2004) at 1892, stating:

The appearance of mere indistinct words in a specification or a claim, even an original claim, does not necessarily satisfy that requirement.

A description of an anti-inflammatory steroid, i.e., a steroid (a generic structural term) described even in terms of its functioning of lessening inflammation of tissues <u>fails to distinguish any steroid from others</u>

having the same activity or function. A description of what a material

does, rather than of what it is, usually does not suffice.... The disclosure must allow one skilled in the art to <u>visualize or recognize</u> the identity of the subject matter purportedly described. (Emphasis added).

Conversely, a description of a chemical genus will usually comprise a recitation of structural features common to the members of the genus, which features constitute a substantial portion of the genus. See <u>Univ. of Calf. V. Eli</u>

<u>Lilly</u>, 43 USPQ 2d 1398, 1406 (Fed. Cir. 1997). This is analogous to enablement of a genus under Section 112, ¶ 1, by showing the enablement of a representative number of species within the genus.

A chemical genus can be adequately described if the disclosure presents a sufficient number of representative species that encompass the genus. If the genus has substantial variance, the disclosure must describe a sufficient number of species to reflect the variation within that genus. See MPEP 2163. The MPEP lists factors that can be used to determine if sufficient evidence of possession has been furnished in the disclosure of the Application. These include the level of skill and knowledge in the art, partial structure, physical and/or chemical properties, functional characteristics alone or coupled with a known or disclosed correlation

between structure and function, and the method of making the claimed invention.

Disclosure of any combination of such identifying characteristics that distinguish the claimed invention from other materials and would lead one of skill in the art to the conclusion that the applicant was in possession of the claimed species is sufficient.

Here, the specification does not provide a reasonably representative disclosure of useful for reducing the level of C-reactive protein (CRP) in an individual subject to a CRP associated inflammatory condition generally, a potentially huge genus inclusive of many different compounds having widely divergent structures and functions.

Applicant is kindly requested to explain the issue. In the present case

Applicant has no possession for the claimed subject matter. Further the

compounds as in claim 1 covers large number of compounds to treat cancer. At the

time invention was filed applicant has no possession of the invention as claimed.

The written description requirement prevents applications from using the amendment process to update the disclosure in their disclosures (claims or specification) during the pendency before the patent office. Otherwise applicants could add new matter to their disclosures and date them back to their original filing date, thus defeating an accurate accounting of the priority of the invention. See 35

Page 9

Art Unit: 1612

USC 132. The function of description requirement is to ensure that the inventor had possession, as of filing date of the application relied on, the specific subject matter claimed by him.

See Genetech, 108 F 3d 1361, 1365 (Fed. Cir. at 1366, 78, 1999).

Possession may be shown in a variety of ways including description of an actual reduction to practice, or by showing that the invention was "ready for patenting" such as by the disclosure of drawings or structural chemical formulas that show that the invention was complete, or by describing distinguishing identifying characteristics sufficient to show that the applicant was in possession of the claimed invention. See, e.g., *Pfaff v. Wells Elecs., Inc.*, 525 U.S. 55, 68, 119 S.Ct. 304, 312, 48 USPQ2d 1641, 1647 (1998); *Eli Lilly*, 119 F.3d at 1568, 43 USPQ2d at 1406; *Amgen, Inc. v. Chugai Pharmaceutical*, 927 F.2d 1200, 1206, 18 USPQ2d 1016, 1021 (Fed. Cir. 1991) (one must define a compound by "whatever characteristics sufficiently distinguish it").

See MPEP 2163.06.

Response to Remarks

Applicant's response filed on 11/18/2008 with traverse the election of species when R is O is hereby acknowledged.. It is unclear that Applicant cites election of claim 2 from group 2 because there are no groups. The request was for

the election one species. Applicant should have elected one compound from claim 2. The basis of the traversal is that Applicants consider that there is no undue burden to examine all the claimed invention. Claims are drawn to a method of reducing the level of C-reactive protein (CRP) in an individual subject to a CRP associated inflammatory condition containing the compound of formula I. In formula I R can be O, S, SO, SO2, a secondary or tertiary amine, a phosphate, a phosphor-ester, or a substituted or unsubstituted methylene group. These compounds include thousands of compounds. Furthermore, n can be 0 to 3, so that the hetero ring will vary depending on n.

Examiner disagrees because the compounds of formula I as in claim 1 contain large number references as was evidenced by the search on STN. It is not possible to go through all the references to determine the patentability of invention. It will be an undue burden on the examiner to examine all the invention as has been claimed. For the same reasons restriction is maintained and is made final.

Communication

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Sabiha Qazi whose telephone number is (571)

272-0622. The examiner can normally be reached on any business day except

Wednesday.

If attempts to reach the examiner by telephone are unsuccessful, the

examiner's supervisor, Krass Frederick can be reached on (571) 272-0580. The

fax phone number for the organization where this application or proceeding is

assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the

Patent Application Information Retrieval (PAIR) system. Status information for

published applications may be obtained from either Private PAIR or Public PAIR.

Status information for unpublished applications is available through Private PAIR

only. For more information about the PAIR system, see http://pair-

direct.uspto.gov. Should you have questions on access to the Private PAIR system,

contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you

would like assistance from a USPTO Customer Service Representative or access to

the automated information system, call 800-786-9199 (IN USA OR CANADA) or

571-272-1000.

/Sabiha Qazi/

Primary Examiner, Art Unit 1612

Application/Control Number: 10/714,152

Page 12

Art Unit: 1612